

K962121 HORIZON/WESTHIN 55-GAug 26, 1996
87 days to decisionK962121 · Product code: **LPL** · Ophthalmic
Source: <https://www.510kdatabase.net/k962121/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lenses, Soft Contact, Daily Wear (LPL)
Date received	May 31, 1996
Decision date	Aug 26, 1996
Days to decision	87 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Westcon Contact Lens Co., Inc.
Location	Grand Junction, CO, US
Contact	MARTIN DALSING
510(k) history	9 submissions · 9 cleared · 1994-2005

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k962121/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 29, 2026